INTERNATIONAL SEARCH REPORT

International application No.

		PCT/JP2	004/015594
A. CLASSIFIC Int.Cl ⁷	ATION OF SUBJECT MATTER C12N5/10, C07K14/745, C12N15,	/12, C07K14/46, C12P21/0)2
	emational Patent Classification (IPC) or to both nationa	I classification and IPC	
B. FIELDS SE.			
Minimum docum Int.C1 ⁷	tentation searched (classification system followed by cl C12N5/10, C07K14/745, C12N15,	assification symbols) /12, C07K14/46, C12P21/0	02
	earched other than minimum documentation to the extension of the extension		•
JSTPlus	S (JOIS), BIOSIS/WPI (DIALOG)	uata tese and, where practicable, scarch te	Ans used)
C. DOCUMEN	ITS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where ap	<u> </u>	Relevant to claim No.
X A	JP 2003-510087 A (Genentech, 18 March, 2003 (18.03.03), Full text & WO 01/23592 A2 & EP & US 6586206 B1	Inc.),	<u>1-22</u> 23
<u>X</u> A	Lin G. et al., Stable cell lines expressing baculovirus P35: resistance to apoptosis and nutrient stress, and increased glycoprotein secretion, In Vitro Cell Dev.Biol.Anim., 2001, Vol.37, No.5, pages 293 to 302		<u>1-22</u> 23
X A	Greenberg C.S. et al., Cleava coagulation factor XIII and a thrombin during in vitro clot Invest., 1985, Vol.75, No.5,	Tibrinogen by ting, J.Clin.	2 <u>3</u> 1-22
× Further do	cuments are listed in the continuation of Box C.	See patent family annex.	
"A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier application or patent but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is		 ater document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention adocument of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone adocument of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art adocument member of the same patent family 	
Date of the actual completion of the international search 02 February, 2005 (02.02.05)		Date of mailing of the international search report 22 February, 2005 (22.02.05)	
Name and mailing address of the ISA/ Japanese Patent Office		Authorized officer Telephone No.	
Form PCT/ISA/21	0 (second sheet) (January 2004)	Telephone No.	

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International application No.
PCT/JP2004/015594

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X A	Roy S.N. et al., Assembly and secretion of recombinant human fibrinogen, J.Biol.Chem., 1991, Vol.266, No.8, pages 4758 to 4763	$\frac{23}{1-22}$ $\frac{23}{1-22}$
X A	Morita T. et al., Purification and properties of prothrombin activator from the venom of Echis carinatus, J.Biochem., 1978, Vol.83, No.2, pages 559 to 570	
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The substance as set forth in claim 23 is specified as "a protein high-produced by using a recombinant animal cell as set forth in any of claims 1 to 17" and, therefore, involves any proteins produced by using these animal cells.

However, fibrinogen, ecarin and factor VIII are exclusively disclosed in the description as proteins produced by using these animal cells. Although the common technical knowledge at the point of the application is taken into consideration, the scope of substances obtained by the screening method cannot be specified. Thus, claim 23 is described in an unclear manner.

Such being the case, the search was made on fibrinogen, ecarin and factor VIII which are specifically presented in the description.